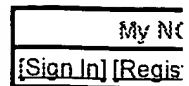
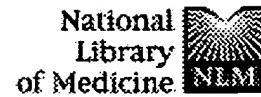


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L7 88746 S PAPILLOMAVIRUS OR PALLOMA VIRUS OR HPV OR HUMAN PAPILLOMAVIRU  
L8 2207 S L7 AND STRAIN#  
L9 219 S L8 AND (PATHOGEN###)  
L10 6 S L9 AND ((SINGLE (3A) PRIMER) OR (ONE (3A) PRIMER) OR PRIMER)  
L11 1 S L10 AND PROBE  
L12 6 S L9 AND PRIMER  
L13 6 S L9 AND PROBE  
L14 6 DUP REM L12 (0 DUPLICATES REMOVED)  
L15 6 DUP REM L14 (0 DUPLICATES REMOVED)  
L16 0 S L9 AND (PNA (10A) AMPLIF?)  
L17 0 S PNA AND L9  
L18 502 S D HIS  
L19 1 S L9 AND LIGASE CHAIN REACTION  
L20 0 S L9 AND ROLLING CIRCLE AMPLIFICATION  
L21 51 S L7 AND (LIGASE CHAIN REACTION)  
L22 21 S L7 AND ROLLING CIRCLE AMPLIFICATION  
L23 88 DUP REM L2 (135 DUPLICATES REMOVED)  
L24 37 DUP REM L21 (14 DUPLICATES REMOVED)  
L25 8 DUP REM L22 (13 DUPLICATES REMOVED)  
L26 2 S L7 AND PNA PROBES

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1: Gynecol Obstet Invest. 1993;36(3):185-8.

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**Human papilloma virus infection and Ki-ras oncogene in paraffin-embedded squamous carcinomas of the cervix.****Falcinelli C, Luzi P, Alberti P, Cosmi EV, Anceschi MM.**

II Institute of Obstetrics and Gynecology, University La Sapienza, Rome, Italy.

42 paraffin-embedded squamous cervical carcinomas were screened for the presence of human papilloma virus (HPV; 6b, 11, 16, 18) and for activation of the Ki-ras oncogene family by polymerase chain reaction. In 72% of cases we found one or more HPV types, but no mutations of the Ki-ras gene (codon 12-1 and 61-1, 61-2 and 61-3) were found. We conclude that mutations of the Ki-ras oncogene, at the positions analyzed, are not likely to be involved in the events leading to cervical carcinogenesis.

PMID: 8244195 [PubMed - indexed for MEDLINE]

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